



US009434050B2

(12) **United States Patent**
Vail

(10) **Patent No.:** **US 9,434,050 B2**

(45) **Date of Patent:** ***Sep. 6, 2016**

(54) **METHODS OF FABRICATING ABRASIVE ELEMENTS USING SP²-CARBON-CONTAINING PARTICLES**

C22C 29/06 (2006.01)

B22F 5/00 (2006.01)

(52) **U.S. Cl.**

CPC . **B24D 3/10** (2013.01); **B22F 7/06** (2013.01);

B82Y 30/00 (2013.01); **C04B 35/52** (2013.01);

C04B 35/645 (2013.01); **C22C 26/00**

(2013.01); **E21B 10/567** (2013.01); **B22F**

2005/001 (2013.01); **C04B 2235/427**

(2013.01); **C04B 2235/5436** (2013.01); **C04B**

2235/5454 (2013.01); **C04B 2235/5472**

(2013.01); **C04B 2235/77** (2013.01); **C22C**

29/06 (2013.01)

(71) Applicant: **US Synthetic Corporation**, Orem, UT (US)

(72) Inventor: **Michael A Vail**, Genola, UT (US)

(73) Assignee: **US SYNTHETIC CORPORATION**, Orem, UT (US)

(*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 523 days.

This patent is subject to a terminal disclaimer.

(58) **Field of Classification Search**

None

See application file for complete search history.

(56)

References Cited

U.S. PATENT DOCUMENTS

3,745,623 A 7/1973 Wentorf, Jr. et al.

4,104,441 A 8/1978 Fedoseev

(Continued)

FOREIGN PATENT DOCUMENTS

EP 0715930 6/1996

EP 07872564 5/2009

(Continued)

OTHER PUBLICATIONS

U.S. Appl. No. 12/245,573, Feb. 6, 2014, Office Action.

(Continued)

Primary Examiner — George Wyszomierski

Assistant Examiner — Ngoclan T Mai

(74) *Attorney, Agent, or Firm* — Dorsey & Whitney LLP

(57)

ABSTRACT

In an embodiment, a method of forming an abrasive element is disclosed. The method includes sintering a powder mixture including a plurality of superabrasive particles, a plurality of sp²-carbon-containing particles, and a metallic constituent to form the abrasive element.

20 Claims, 4 Drawing Sheets

(21) Appl. No.: **13/912,433**

(22) Filed: **Jun. 7, 2013**

(65) **Prior Publication Data**

US 2013/0269263 A1 Oct. 17, 2013

Related U.S. Application Data

(63) Continuation of application No. 13/533,026, filed on Jun. 26, 2012, now Pat. No. 8,936,117, which is a continuation of application No. 13/116,511, filed on May 26, 2011, now Pat. No. 8,246,701, which is a continuation of application No. 12/394,594, filed on Feb. 27, 2009, now Pat. No. 7,972,397, which is a continuation of application No. 11/496,905, filed on Jul. 31, 2006, now Pat. No. 7,516,804.

(51) **Int. Cl.**

B22F 7/06 (2006.01)

B24D 3/10 (2006.01)

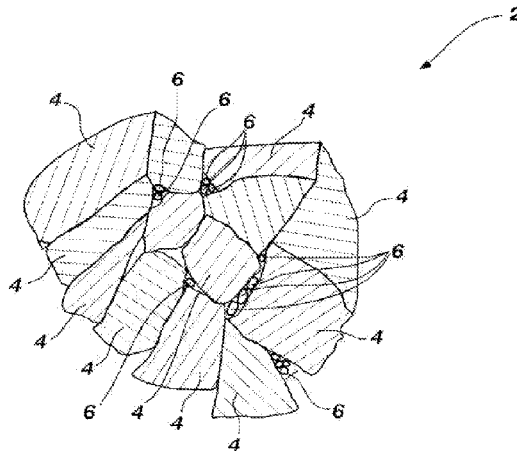
E21B 10/567 (2006.01)

C04B 35/52 (2006.01)

C04B 35/645 (2006.01)

C22C 26/00 (2006.01)

B82Y 30/00 (2011.01)



(56)

References Cited

U.S. PATENT DOCUMENTS

4,246,005	A	1/1981	Ishizuka	
4,268,276	A	5/1981	Bovenkerk	
4,274,900	A	6/1981	Mueller et al.	
4,410,054	A	10/1983	Nagel et al.	
4,425,315	A	1/1984	Tsuji et al.	
4,468,138	A	8/1984	Nagel	
4,560,014	A	12/1985	Geczy	
4,636,253	A	1/1987	Nakai et al.	
4,738,322	A	4/1988	Hall et al.	
4,797,241	A	1/1989	Peterson et al.	
4,811,801	A	3/1989	Salesky et al.	
4,850,523	A *	7/1989	Slutz	C04B 37/006 228/103
4,913,247	A	4/1990	Jones	
5,016,718	A	5/1991	Tandberg	
5,087,435	A	2/1992	Potter et al.	
5,092,687	A	3/1992	Hall	
5,120,327	A	6/1992	Dennis	
5,128,080	A	7/1992	Jurewicz	
5,135,061	A	8/1992	Newton, Jr.	
5,154,245	A	10/1992	Waldenstrom et al.	
5,209,916	A	5/1993	Gruen	
5,328,676	A	7/1994	Gruen	
5,364,192	A	11/1994	Damm et al.	
5,368,398	A	11/1994	Damm et al.	
5,370,855	A	12/1994	Gruen	
5,449,491	A	9/1995	Job	
5,460,233	A	10/1995	Meany et al.	
5,462,776	A	10/1995	Gruen	
5,467,836	A	11/1995	Grimes et al.	
5,480,233	A	1/1996	Cunningham	
5,544,713	A	8/1996	Dennis	
5,620,512	A	4/1997	Gruen et al.	
5,759,216	A	6/1998	Kanada et al.	
5,855,996	A *	1/1999	Corrigan	B01J 3/062 428/212
5,900,225	A	5/1999	Mistry et al.	
5,954,147	A	9/1999	Overstreet et al.	
6,214,079	B1	4/2001	Kear et al.	
6,398,815	B1	6/2002	Pope et al.	
6,544,308	B2	4/2003	Griffin et al.	
6,783,745	B1	8/2004	Voronov et al.	
6,793,681	B1	9/2004	Pope et al.	
6,800,095	B1	10/2004	Pope et al.	
6,883,623	B2	4/2005	McCormick et al.	
7,048,081	B2	5/2006	Smith et al.	
7,060,641	B2	6/2006	Qian et al.	
7,070,635	B2	7/2006	Frushour	
7,350,599	B2	4/2008	Lockwood et al.	
7,516,804	B2	4/2009	Vail	
7,517,588	B2	4/2009	Frushour	
7,569,176	B2	8/2009	Pope et al.	
7,635,035	B1	12/2009	Bertagnolli	
7,841,428	B2	11/2010	Bertagnolli	
7,842,111	B1	11/2010	Sani	
7,972,397	B2	7/2011	Vail	
8,246,701	B2	8/2012	Vail	
2004/0011433	A1	1/2004	Shiozaki et al.	
2004/0057896	A1	3/2004	Kronholm et al.	
2005/0002851	A1	1/2005	McElrath et al.	
2005/0019114	A1	1/2005	Sung	
2005/0133277	A1	6/2005	Dixon	
2005/0186104	A1	8/2005	Kear et al.	
2005/0227590	A1	10/2005	Sung	
2006/0016127	A1	1/2006	Sung	
2006/0042172	A1	3/2006	Sung	
2006/0060390	A1	3/2006	Eyre	
2006/0060392	A1	3/2006	Eyre	
2006/0086540	A1	4/2006	Griffin et al.	
2006/0147644	A1	7/2006	Fujimura et al.	
2006/0157285	A1	7/2006	Cannon et al.	
2006/0266559	A1	11/2006	Keshavan et al.	
2007/0056778	A1	3/2007	Webb et al.	
2007/0144790	A1	6/2007	Fang et al.	
2007/0187153	A1	8/2007	Bertagnolli	

2007/0234646	A1	10/2007	Can et al.
2007/0272448	A1	11/2007	Griffo
2008/0085407	A1	4/2008	Cooley et al.
2008/0115424	A1	5/2008	Can et al.
2009/0152015	A1	6/2009	Sani et al.
2009/0178345	A1	7/2009	Russell et al.
2010/0104874	A1	4/2010	Yong et al.
2012/0267174	A1	10/2012	Vail

FOREIGN PATENT DOCUMENTS

EP	07872564	6/2011
FR	2684090	5/1993
RU	2131763	6/1999
WO	WO 99/10274	3/1999
WO	WO 2008/094190	8/2008

OTHER PUBLICATIONS

U.S. Appl. No. 12/912,273, Oct. 25, 2013, Notice of Allowance.
U.S. Appl. No. 12/912,273, Mar. 3, 2014, Notice of Allowance.
U.S. Appl. No. 13/533,026, Dec. 6, 2013, Office Action.
U.S. Appl. No. 12/912,273, Apr. 8, 2014, Notice of Allowance.
U.S. Appl. No. 12/912,273, May 7, 2014, Issue Notification.
U.S. Appl. No. 12/245,573, Oct. 3, 2008, Sani.
U.S. Appl. No. 12/909,716, Oct. 21, 2010, Bertagnolli.
U.S. Appl. No. 12/912,273, Oct. 26, 2010, Sani.
Rhede, D. et al. "Diamond powder-contg. Moulding prodn. Using expanded graphite addn. improved mixability and pressability", Derwent Abstract of DD26888, Jun. 1989.
Akaishi, Minoru, Yoichiro Sato, Nobuo Setaka, Masayuki Tsutsumi, Toshikazu Ohsawa and Osamu Fukunaga, "Effect of Additive Graphite on Sintering of Diamond", Cermaic Bulletin, vol. 62, No. 6 (1983) pp. 689-694.
Asbury Carbons, "Amorphous Graphite", <http://asbury.com/Amorphous-Graphite.html> (Mar. 28, 2008) (May 16, 2012) (obtained via Internet Archive Wayback Machine).
Davidenko, V.M., S.V. Kidalov, F.M. Shakhov, M.A. Yagovkina, V.A. Yashin, V.Ya. Vul, "Fullerenes as a co-catalyst for high pressure-high temperature synthesis of diamonds", Diamond and Related Materials 13 (2004) pp. 2203-2206.
Kidalov, S.V., V.I. Sokolov, F.M. Shakhov and A. Ya. Vul', "Mechanism of the Catalytic Effect of Fullerenes on the Graphite-Diamond Phase Transition at High Temperature and Pressure", Doklady Physical Chemistry, vol. 404, Part 1, (2005) pp. 179-181.
Vul, A. Ya, V.M. Davidenko, S.V. Kidalov, S.S. Ordan'yan and V.A. Yashin, "Fullereness Catalyze the Graphite-Diamond Phase Transition", Technical Physics Letters, vol. 27, No. 5, 2001, pp. 384-386.
Vul, A. Ya, S.V. Kidalov, V.M. Davidenko, V.A. Yashin, S.S. Ordanyan, "Fullereness as a co-catalyst for HPHT Synthesis of Diamond", Proceedings of the Sixth Applied Diamond Conference/ Second (2001) pp. 237-238.
Yushin, G.N. et al., "Effect of sintering in structure of nanodiamond," Diamond and Related Materials, Aug. 10, 2005, vol. 14, pp. 1721-1729.
International Search Report and Written Opinion from International Application No. PCT/US2007/016322 dated Oct. 17, 2008.
U.S. Appl. No. 11/496,905, Apr. 17, 2008, Office Action.
U.S. Appl. No. 11/496,905, Oct. 30, 2008, Office Action.
U.S. Appl. No. 11/496,905, Jan. 15, 2009, Notice of Allowance.
U.S. Appl. No. 11/496,905, Mar. 25, 2009, Issue Notification.
U.S. Appl. No. 11/351,564, May 27, 2008, Restriction Requirement.
U.S. Appl. No. 11/351,564, Sep. 17, 2008, Office Action.
U.S. Appl. No. 11/351,564, Mar. 4, 2009, Office Action.
U.S. Appl. No. 11/351,564, Jun. 24, 2009, Office Action.
U.S. Appl. No. 11/351,564, Dec. 30, 2009, Notice of Allowance.
U.S. Appl. No. 11/351,564, Apr. 15, 2010, Notice of Allowance.
U.S. Appl. No. 11/351,564, Jul. 21, 2010, Notice of Allowance.
U.S. Appl. No. 11/351,564, Nov. 10, 2010, Issue Notification.
U.S. Appl. No. 12/111,769, Dec. 23, 2009, Restriction Requirement.
U.S. Appl. No. 12/111,769, Mar. 11, 2010, Office Action.
U.S. Appl. No. 12/111,769, Jul. 28, 2010, Notice of Allowance.
U.S. Appl. No. 12/111,769, Aug. 20, 2010, Notice of Allowance.
U.S. Appl. No. 12/111,769, Nov. 10, 2010, Issue Notification.

(56)

References Cited

OTHER PUBLICATIONS

U.S. Appl. No. 12/394,594, Oct. 1, 2010, Restriction Requirement.
U.S. Appl. No. 12/394,594, Dec. 22, 2010, Office Action.
U.S. Appl. No. 12/394,594, Mar. 31, 2011, Notice of Allowance.
U.S. Appl. No. 12/394,594, Jun. 15, 2011, Issue Notification.
U.S. Appl. No. 12/245,573, Jun. 17, 2011, Restriction Requirement.
U.S. Appl. No. 12/245,573, Oct. 26, 2011, Office Action.
U.S. Appl. No. 12/245,573, Jun. 11, 2012, Office Action.

U.S. Appl. No. 12/909,716, Feb. 14, 2012, Office Action.
U.S. Appl. No. 13/116,511, Dec. 9, 2011, Office Action.
U.S. Appl. No. 13/116,511, Mar. 30, 2012, Notice of Allowance.
U.S. Appl. No. 14/569,039, filed Dec. 12, 2014, Sani.
U.S. Appl. No. 12/245,573, Sep. 10, 2014, Notice of Allowance.
U.S. Appl. No. 12/245,573, Mar. 4, 2015, Issue Notification.
U.S. Appl. No. 13/533,026, Sep. 15, 2014, Notice of Allowance.
U.S. Appl. No. 13/533,026, Dec. 30, 2014, Issue Notification.
U.S. Appl. No. 14/569,039, Jul. 26, 2016, Restriction Requirement.

* cited by examiner

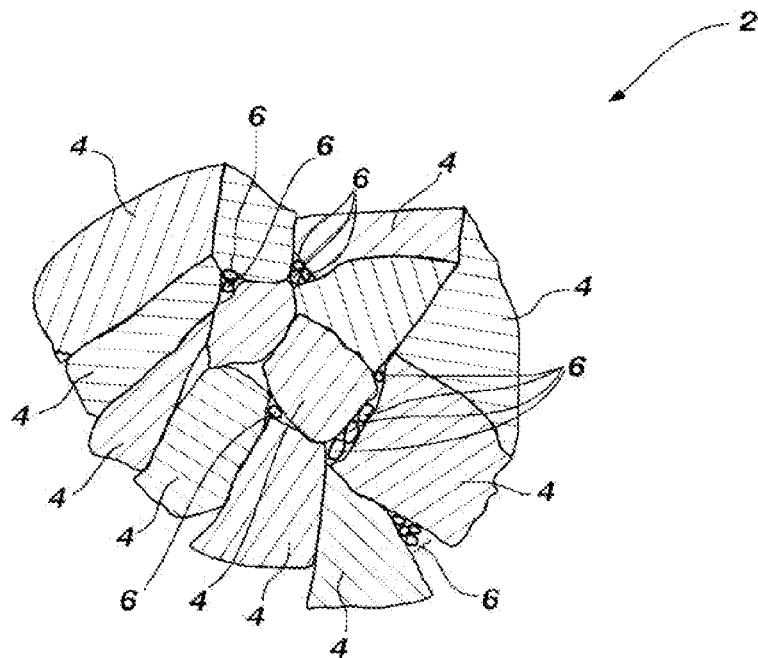


FIG. 1

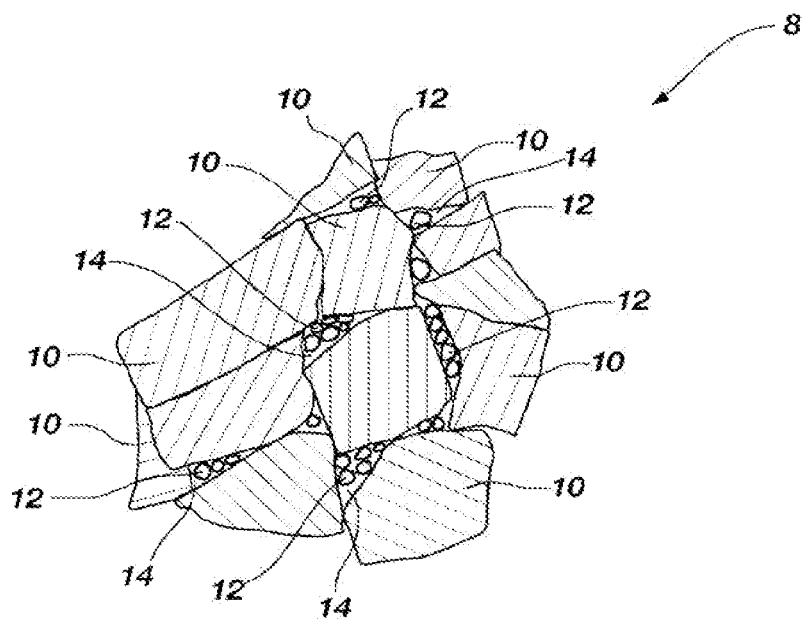


FIG. 2

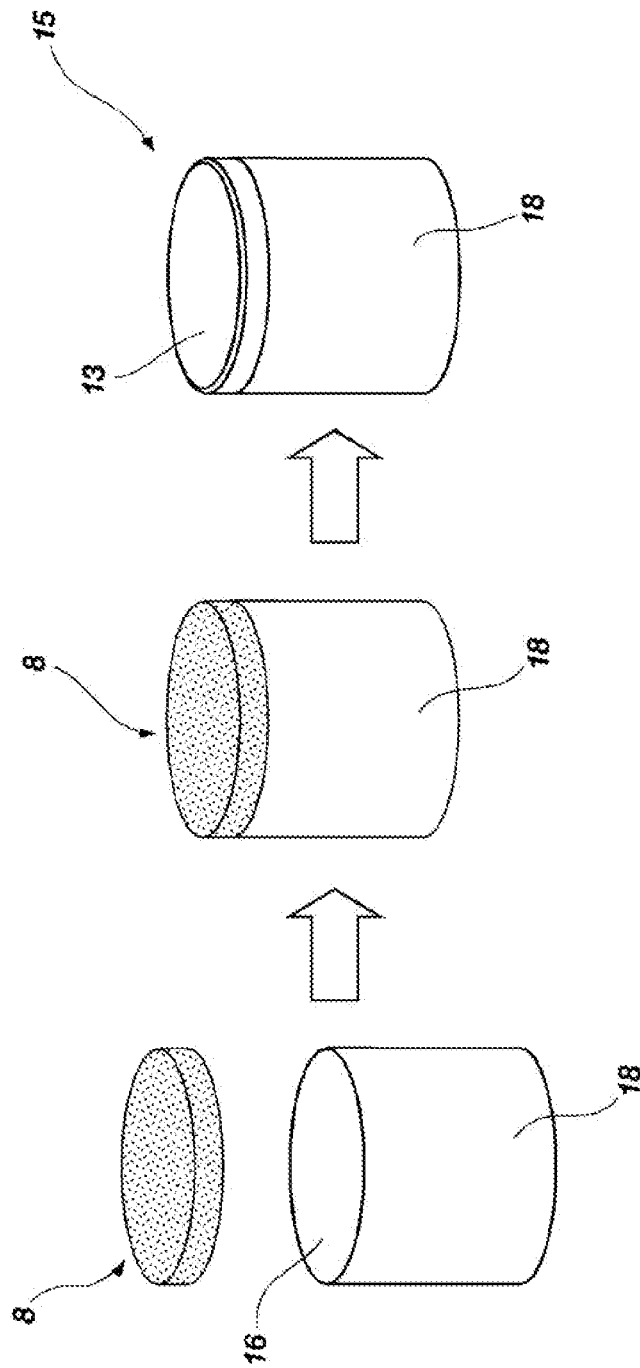


FIG. 3

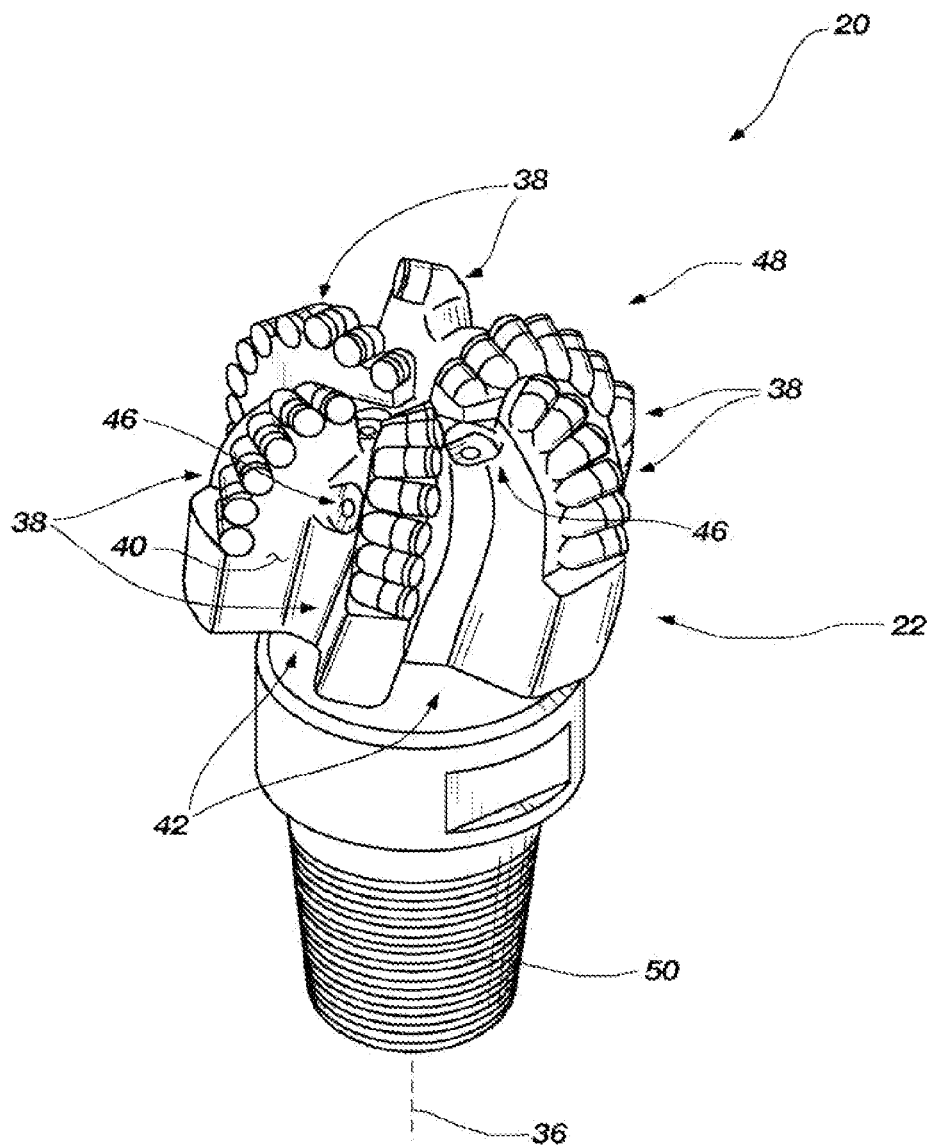


FIG. 4

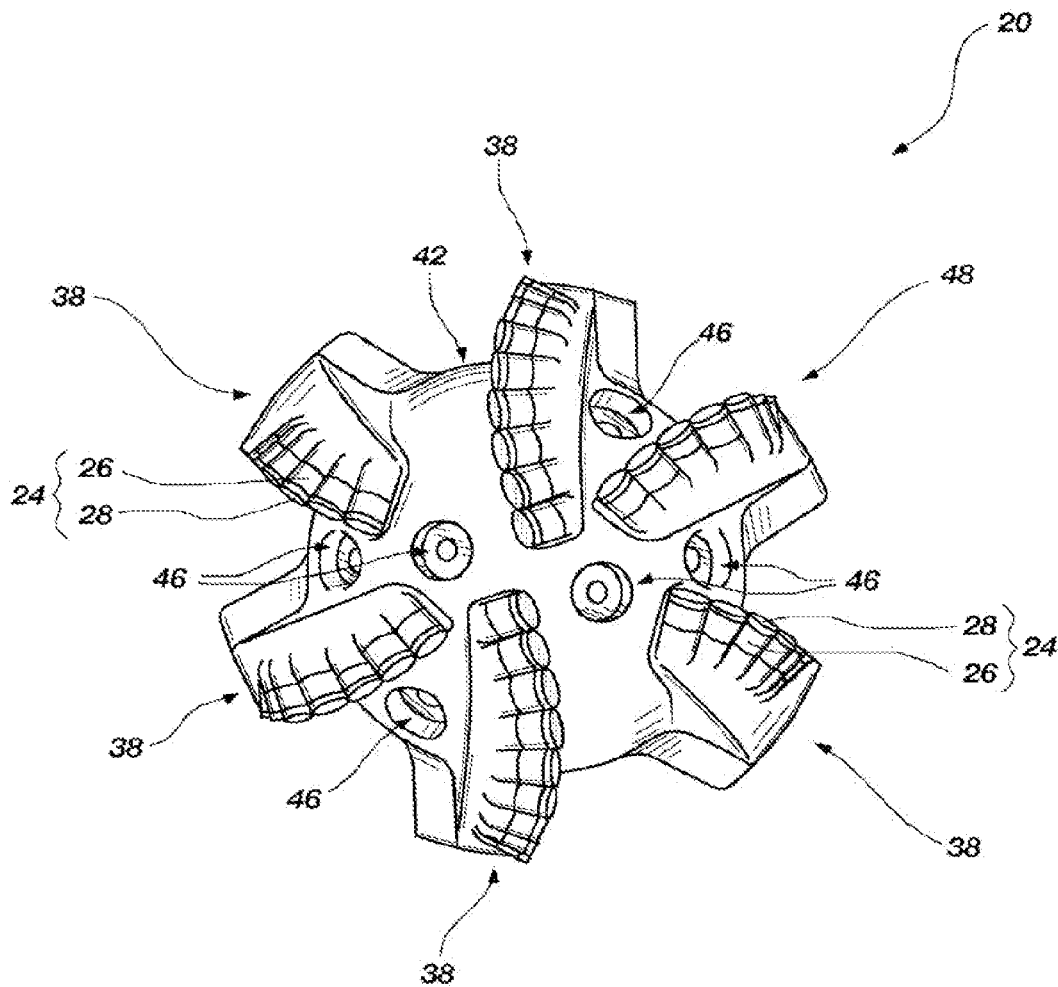


FIG. 5

1

METHODS OF FABRICATING ABRASIVE ELEMENTS USING SP2-CARBON-CONTAINING PARTICLES

CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a continuation of application Ser. No. 13/533,026 filed on 26 Jun. 2012, which is a continuation of application Ser. No. 13/116,511 filed on 26 May 2011 (now U.S. Pat. No. 8,246,701 issued on 21 Aug. 2012), which is a continuation of application Ser. No. 12/394,594 filed on 27 Feb. 2009 (now U.S. Pat. No. 7,972,397 issued on 5 Jul. 2011), which is a continuation of application Ser. No. 11/496,905 filed on 31 Jul. 2006 (now U.S. Pat. No. 7,516,804 issued on 14 Apr. 2009). Each of the foregoing applications is incorporated, in its entirety, by this reference.

TECHNICAL FIELD

The present invention relates to superabrasive materials, apparatuses, and methods of manufacturing same, such as polycrystalline diamond (PCD) elements and applications utilizing such elements in drilling tools (e.g., inserts, cutting elements, gage trimmers, etc.), machining equipment, bearing apparatuses, wire drawing machinery, and other apparatuses.

BACKGROUND

Polycrystalline diamond compacts ("PDCs"), such as cutting elements used in rock bits or other cutting tools, typically include a cemented tungsten carbide substrate having a layer of superabrasive PCD (also commonly referred to as a diamond table) bonded to a surface of the substrate using an ultra-high pressure, ultra-high temperature ("HPHT") process. Sometimes, the substrate may be brazed or otherwise joined to an attachment member such as a stud or to a cylindrical backing, if desired. A stud carrying a PDC may be used as a subterranean cutting element when mounted to a drill bit by press-fitting, brazing, or otherwise locking the stud into a receptacle formed in the subterranean drill bit or by brazing the cutting element directly into a preformed pocket, socket, or other receptacle formed in the subterranean drill bit. For example, cutter pockets may be formed in the face of a bit formed of cemented tungsten carbide. Generally, a rotary drill bit may include a plurality of PCD superabrasive cutting elements affixed to the drill bit body.

Conventional PDCs are normally fabricated by placing a cemented carbide substrate into a container or cartridge with a volume of diamond crystals or particles positioned on a surface of the substrate. A number of such cartridges may be typically loaded into an ultra-high pressure press. The substrates and volume of diamond particles are then processed under HPHT conditions in the presence of a catalyst material that causes the diamond grains to form and to bond to one another to form a matrix of bonded diamond grains defining a diamond table. The catalyst material is often a solvent catalyst, such as cobalt, nickel, or iron that is used for facilitating the intergrowth of the diamond grains. In one process, a constituent of the substrate, such as cobalt from a cobalt-cemented tungsten carbide substrate, becomes liquid and sweeps from the region adjacent to the volume of diamond grains and into interstitial regions between the diamond grains during the HPHT process. The cobalt acts as a catalyst to facilitate the intergrowth process between the

2

diamond grains, which results in bonds between adjacent diamond grains. Often, a solvent catalyst may be mixed with the diamond particles prior to subjecting the diamond particles and the substrate to the HPHT process.

As known in the art, the solvent catalyst may dissolve carbon from the diamond particles or portions of the diamond particles that graphitize due to the high temperatures being used. The solubility of the stable diamond phase in the solvent catalyst is lower than that of the metastable graphite under HPHT conditions. As a result of this solubility difference, the undersaturated graphite tends to dissolve into solvent catalyst and the supersaturated diamond tends to deposit onto existing diamond grains to form diamond-to-diamond bonds. Accordingly, diamond grains become mutually bonded to form a matrix of PCD with interstitial regions between the bonded diamond grains being occupied by the solvent catalyst.

However, the presence of the solvent catalyst in the diamond table can lead to a diamond table that may be thermally damaged at elevated temperatures. For example, the difference in thermal expansion coefficient between the diamond grains and the solvent catalyst is believed to lead to chipping or cracking in the PDC during drilling or cutting operations, which consequently can degrade the mechanical properties of the PDC or cause failure. Additionally, it is believed that some of the diamond grains can undergo a chemical breakdown or back-conversion with the solvent catalyst. Of course, at extremely high temperatures, diamond may transform to carbon monoxide, carbon dioxide, graphite, or combinations thereof degrading the mechanical properties of the PDC.

Therefore, there is still a need for a superabrasive material (e.g., PCD) exhibiting superior mechanical and/or thermal properties (e.g., an increased amount of bonding between superabrasive grains).

SUMMARY

The present invention is directed to superabrasive materials and elements, such as PCD elements, methods of fabricating superabrasive elements, and applications utilizing such elements. One aspect of the present invention is directed to a superabrasive element comprising a mass of polycrystalline diamond including ultra-dispersed diamond grain structures present in an amount greater than zero weight percent and less than about 75 weight percent of the mass of polycrystalline diamond. In one embodiment, the ultra-dispersed diamond grain structures may be present in an amount greater than zero weight percent and less than about 45 weight percent of the mass of polycrystalline diamond.

An additional aspect of the present invention is directed to a method of fabricating a superabrasive element. A mixture including a first type of superabrasive particles comprising ultra-dispersed diamond particles and a second type of superabrasive particles is provided. The ultra-dispersed diamond particles are present in an amount greater than zero weight percent and less than about 75 weight percent of the mixture. The mixture is sintered by application of heat and pressure in an amount sufficient to form the superhard element. In one embodiment, the ultra-dispersed diamond particles may be present in an amount greater than zero weight percent and less than about 45 weight percent of the mixture.

Another aspect of the present invention is directed to a PDC. The PDC includes a substrate and a superabrasive table bonded to the substrate. The superabrasive table

includes a mass of polycrystalline diamond comprising ultra-dispersed diamond grain structures present in an amount greater than zero weight percent and less than about 75 weight percent of the mass of polycrystalline diamond. In one embodiment, the ultra-dispersed diamond grain structures may be present in an amount greater than zero weight percent and less than about 45 weight percent of the mass of polycrystalline diamond.

A further aspect of the present invention is directed to a drill bit. The drill bit includes a bit body adapted to engage a subterranean formation during drilling. At least one superabrasive cutting element is affixed to the bit body. The at least one superabrasive cutting element includes a mass of polycrystalline diamond comprising ultra-dispersed diamond grain structures present in an amount greater than zero weight percent and less than about 75 weight percent of the mass of polycrystalline diamond. In one embodiment, the ultra-dispersed diamond grain structures may be present in an amount greater than zero weight percent and less than about 45 weight percent of the mass of polycrystalline diamond.

Yet a further aspect of the present invention is directed to various different apparatuses, such as drilling tools, machining equipment, bearing apparatuses, wire drawing machinery, and other apparatuses that employ the inventive superabrasive elements disclosed herein.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a partial, schematic microstructural representation of a sintered superabrasive element in accordance with one embodiment of the present invention.

FIG. 2 is a partial, schematic microstructural representation of a mixture of a first type of superabrasive particles formed of relatively coarse superabrasive particles and a second type of superabrasive particles formed of ultra-dispersed diamond particles that are used to fabricate the superabrasive element of FIG. 1 in accordance with one embodiment of a method of the present invention.

FIG. 3 is a schematic diagram illustrating a method for fabricating a superabrasive element on a substrate in accordance with one embodiment of the present invention.

FIG. 4 is an isometric view of one embodiment of a rotary drill bit including at least one cutting element comprising a superabrasive element fabricated and structured in accordance with embodiments of the present invention; and

FIG. 5 shows a top elevation view of the rotary drill bit of FIG. 4.

DETAILED DESCRIPTION OF VARIOUS EMBODIMENTS

The present invention is directed to superabrasive (e.g., PCD) elements, methods of fabricating such elements, and applications utilizing such elements. The superabrasive elements disclosed herein may be used in a variety of applications such as drilling tools (e.g., compacts, inserts, cutting elements, gage trimmers, etc.), machining equipment, bearing apparatuses, wire drawing machinery, and other apparatuses. The present invention relates generally to structures comprising at least one superabrasive material (e.g., diamond, boron nitride, silicon carbide, mixtures of the foregoing, or any material exhibiting a hardness exceeding a hardness of tungsten carbide) and methods of manufacturing such structures. More particularly, the present invention relates to a sintered superabrasive mass or volume and methods of manufacturing such a material. As used herein,

“superabrasive,” means a material exhibiting a hardness exceeding a hardness of tungsten carbide. For example, ultra-dispersed diamond particles may be employed for enhancing polycrystalline diamond. In another example, ultra-dispersed diamond particles may be employed for enhancing a superabrasive material as disclosed in U.S. Pat. No. 7,060,641, the disclosure of which is incorporated herein, in its entirety, by this reference.

Many specific details of certain embodiments of the present invention are set forth in the following description and in FIGS. 1 through 5 in order to provide a thorough understanding of such embodiments. While the embodiments described below reference PCD or PDC material, one skilled in the art will understand that the present invention may have additional embodiments, or that the present invention may be practiced without several of the details described in the following description. In the figures and description below, like or similar reference numerals are used to represent like or similar elements.

Various embodiments of the present invention are directed to sintered superabrasive elements that include a mass of sintered superabrasive particles comprising ultra-dispersed diamond grain structures present in an amount greater than zero weight percent and less than about 75 weight percent of the mass of sintered superabrasive particles. As used herein, “ultra-dispersed diamond particles,” also commonly referred to in the art as nanocrystalline diamond particles, means particles each comprising a polycrystalline diamond core surrounded by a metastable carbon shell. Such ultra-dispersed diamond particles may exhibit a particle size of about 1 nm to about 50 nm and more typically, of about 2 nm to about 20 nm, and agglomerates of ultra-dispersed diamond particles may be between about 2 nm to about 200 nm. Ultra-dispersed diamond may be formed by detonating trinitrotoluene explosives in a chamber and subsequent purification to extract diamond particles or agglomerates of diamond particles with the diamond particles generally comprising a polycrystalline diamond core surrounded by a metastable shell, which includes amorphous carbon, carbon onion (i.e., closed shell sp^2 nanocarbons), or both. Ultra-dispersed diamond particles are commercially available from ALIT Inc. of Kiev, Ukraine. The present invention contemplates that as a result of, at least in part, utilizing ultra-dispersed diamond particles in the fabrication process for forming a superabrasive material, the sintered superabrasive material (e.g., PCD) may exhibit enhanced bonding between superabrasive (e.g., diamond) grains and, consequently, increased density, thermal stability, and/or other enhanced physical and/or mechanical properties.

In general, as used herein, the term “particle” or variants thereof, refers to the powder employed prior to sintering a superabrasive material. Further, the term “grain” or the phrase “grain structures,” or variants thereof, refers to discernable superabrasive regions subsequent to sintering, as known and as determined in the art. The phrase “ultra-dispersed diamond grain structure,” as used herein, refers to an ultra-dispersed diamond particle that has been exposed to an HPHT process, without regard to whether the metastable shell exists subsequent to sintering. As used herein, the phrase “HPHT process” means to generate a pressure of at least about 40 kilobar and a temperature of at least about 1000° C.

FIG. 1 shows a partial, schematic microstructural representation of a sintered superabrasive element 2 in accordance with one embodiment of the present invention. The superabrasive element 2 includes coarse diamond grains 4 (represented by the large cross-hatched regions) that are

5

bonded to each other to define grain boundaries. The interstitial regions between adjacent coarse diamond grains **4** are occupied by relatively fine ultra-dispersed diamond grain structures **6** that are bonded to the coarse diamond grains **4** and, depending upon the size of the interstitial regions, other ultra-dispersed diamond grain structures **6**. Although not illustrated in FIG. 1, it should be understood that the ultra-dispersed diamond grain structures **6** exhibit a polycrystalline grain structure.

The ultra-dispersed diamond grain structures **6** are present in the superabrasive element **2** in an amount greater than zero weight percent and less than about 75 weight percent, and in a more specific embodiment the ultra-dispersed diamond grain structures **6** are present in an amount less than amount 45 weight percent of the superabrasive element **2**. In another embodiment, the ultra-dispersed diamond grain structures **6** may be present in an amount greater than zero weight percent to about 5 weight percent, with the balance at least substantially being the coarse diamond grains **4**. In another embodiment, the ultra-dispersed diamond grain structures **6** may be present in an amount greater than zero weight percent to about 4 weight percent, with the balance at least substantially being the coarse diamond grains **4**. In another embodiment, the ultra-dispersed diamond grain structures **6** are present in the superabrasive element **2** in an amount from about 0.1 weight percent to about 3 weight percent, and more specifically in an amount from about 0.1 weight percent to about 2 weight percent, with the balance at least substantially being the coarse diamond grains **4**. In another specific detailed embodiment, the ultra-dispersed diamond grain structures **6** are present in the superabrasive element **2** in an amount from about 0.1 weight percent to about 1.5 weight percent, and more specifically in an amount from about 0.1 weight percent to about 1 weight percent, with the balance at least substantially being the coarse diamond grains **4**. In yet a further detailed embodiment, the ultra-dispersed diamond grain structures **6** are present in the superabrasive element **2** in an amount from about 0.1 weight percent to about 0.5 weight percent, and more specifically in an amount from about 0.1 weight percent to about 0.25 weight percent, with the balance at least substantially being the coarse diamond grains **4**. In yet another detailed embodiment, the ultra-dispersed diamond grain structures **6** are present in the superabrasive element **2** in an amount greater than zero to about 0.1 weight percent with the balance at least substantially being the coarse diamond grains **4**.

The coarse diamond grains **4** exhibit an average grain size (i.e., sintered) that is greater than the average size of the ultra-dispersed diamond grain structures **6**. In a more detailed embodiment, the average grain size of the coarse diamond grains **4** is about 0.5 μm to about 150 μm and the average size of the ultra-dispersed diamond grain structures **6** is about 1 nm to about 50 nm, and, more specifically, about 4 nm to about 30 nm. In another embodiment, the coarse diamond grains **4**, the ultra-dispersed diamond grain structures **6**, or both may exhibit a bimodal or greater distribution of grain/particle sizes. It should be understood that any of the above ranges of various grain sizes and weight percentages for the coarse diamond grains **4** and the ultra-dispersed diamond grain structures **6** may be combined.

The sintered superabrasive element **2** may exhibit a density of at least about 92 percent of theoretical density and may exceed at least about 98 to at least about 99 percent of theoretical density. The superabrasive element **2** further may exhibit an increased extent of diamond-to-diamond bonding between the coarse diamond grains **4**, the ultra-dispersed diamond grain structures **6**, or both. The superabrasive

6

element **2**, formed of substantially only the coarse diamond grains **4** and ultra-dispersed diamond grain structures **6**, may be more thermally stable than a comparative prior art PCD superabrasive element including a solvent catalyst because it does not suffer from deleterious problems commonly experienced when a solvent catalyst is used. However, in some embodiments, a solvent catalyst (e.g., cobalt, nickel, or iron) may also be employed, in at least small amounts, to promote intergrowth of the diamond grains **4** and the ultra-dispersed diamond grain structures **6** and will be present in the interstitial regions between the coarse diamond grains **4**, the ultra-dispersed diamond grain structures **6**, or both. However, such an embodiment may have a comparatively reduced thermal stability.

However, the present invention also contemplates that if the superabrasive element **2** is formed with both ultra-dispersed diamond particles and a catalyst, at least a portion of the catalyst may be subsequently (i.e., after sintering) removed from the PCD element. For example, subsequent to sintering, a catalyst material may be at least partially removed (e.g., by acid-leaching, by a method as otherwise known in the art, or by any suitable method) from at least a portion of a mass of polycrystalline diamond (e.g., a table formed upon a substrate). Catalyst removal may be substantially complete to a selected depth from an exterior surface of the polycrystalline diamond table, if desired, without limitation. Thus, at least a portion of a polycrystalline diamond mass may be substantially free of a catalyst employed during sintering of the polycrystalline diamond mass. As known in the art, at least partial catalyst removal may provide a polycrystalline diamond material with increased thermal stability, which may also beneficially affect the wear resistance of the polycrystalline diamond material.

The superabrasive elements disclosed herein may be fabricated in accordance with various embodiments. For example, referring to FIG. 2, in one embodiment of a method of the present invention, a mixture **8** is formed by mixing at least one type of superabrasive (e.g., diamond) particle formed of relatively coarse superabrasive particles **10** with a selected amount of relatively fine ultra-dispersed diamond particles **12**. The ultra-dispersed diamond particles **12** may be thoroughly cleaned prior to forming the mixture **8** by heating the ultra-dispersed diamond particles **12** in a vacuum furnace for a sufficient time and temperature. The size of the coarse superabrasive particles **10** and the ultra-dispersed diamond particles **12** may be selected so that most of the interstitial regions **14** between the coarse superabrasive particles **10** are occupied by one or more of the ultra-dispersed diamond particles **12**. In one embodiment, the superabrasive particles **10** may comprise diamond (e.g., diamond powder or grit). As known in the art, the mixture **8** may be placed in a pressure transmitting medium such as a refractory metal can, graphite structure, pyrophyllite or other pressure transmitting structures, or other containers or supporting elements. The pressure transmitting medium, including the mixture **8**, is subjected to an HPHT sintering process using an ultra-high pressure press at process conditions of, for example, a pressure of at least about 40 kilobar to at least about 70 kilobar and a temperature of at least about 1000° C. to at least about 1600° C. for a time sufficient to consolidate the mixture and form a coherent mass of bonded diamond shown in FIG. 1 as the superabrasive element **2**.

During the HPHT sintering process, the coarse superabrasive particles **10** of the mixture **8** that are in contact with each other will become bonded to each other. As previously

discussed, most of the ultra-dispersed diamond particles **12** may occupy the interstitial regions **14** between the coarse superabrasive particles **10**. Although the physical phenomenon is not entirely understood, it is currently believed by the inventor that the amorphous carbon, the carbon onion, or both in the ultra-dispersed diamond particles **12** undergo a phase transformation to diamond and bond to the coarse superabrasive particles **10**. Thus, it is believed that the amorphous carbon and/or the carbon onions present in the ultra-dispersed diamond particles **12** are not stable phases at the pressures and temperatures used in the HPHT process. Of course, as shown in FIG. 2, the interstitial regions **14** may include more than one of the ultra-dispersed diamond particles **12** disposed therein. Consequently, as depicted in FIG. 1, the interstitial regions between the coarse diamond grains **4** may include a plurality of ultra-dispersed diamond grain structures **6** bonded to each other. As discussed above, by forming the superabrasive element **2** using the coarse superabrasive particles **10** and the metastable ultra-dispersed diamond particles **12** selected to fit into the interstitial regions **14**, the resulting sintered superabrasive element **2** (FIG. 1) may exhibit enhanced bonding between diamond grains and may exhibit a density close to theoretical density, increased thermal stability, or other enhanced physical and/or mechanical properties.

Prior to sintering, in one embodiment, the coarse superabrasive particles **10** may exhibit an average size of about 0.5 μm to about 150 μm and the ultra-dispersed diamond particles **12** may exhibit an average size of about 1 nm to about 50 nm, with agglomerates of the ultra-dispersed diamond particles **12** being about 2 nm to about 200 nm.

In accordance with one detailed embodiment, mixture **8** may comprise ultra-dispersed diamond particles **12** in an amount greater than zero weight percent and less than about 75 weight percent, and more specifically in an amount less than about 45 weight percent, with the balance being the coarse superabrasive particles **10**. In another embodiment, the mixture **8** may comprise ultra-dispersed diamond particles **12** in an amount greater than zero weight percent and less than about 5 weight percent, and more specifically in an amount greater than zero weight percent to about 4 weight percent, with the balance being the coarse superabrasive particles **10**. In a more detailed embodiment, the ultra-dispersed diamond particles **12** are present in the mixture **8** in an amount from about 0.1 weight percent to about 3 weight percent, and more specifically in an amount from about 0.1 weight percent to about 2 weight percent, with the balance being the coarse superabrasive particles **10**. In another more detailed embodiment, the ultra-dispersed diamond particles **12** are present in the mixture **8** in an amount from about 0.1 weight percent to about 1.5 weight percent, and more specifically in an amount from about 0.1 weight percent to about 1 weight percent, with the balance being the coarse diamond particles **10**. In yet a more specific detailed embodiment, the ultra-dispersed diamond particles **12** are present in the mixture **8** in an amount from about 0.1 weight percent to about 0.5 weight percent, and more specifically in an amount from about 0.1 weight percent to about 0.25 weight percent, with the balance being the coarse superabrasive particles **10**. In yet another more specific detailed embodiment, the ultra-dispersed diamond particles **12** are present in the mixture **8** in an amount greater than zero weight percent to about 0.1 weight percent with the balance being the coarse superabrasive particles **10**.

In another more detailed embodiment, the coarse superabrasive particles **10** of the mixture **8** may exhibit a

bimodal or greater distribution of particle sizes. For example, the mixture **8** may include ultra-dispersed diamond particles **12** in an amount of about 0.1 weight percent to less than about 5 weight percent of the mixture **8** with the balance being coarse superabrasive particles **10**. The formulation of the coarse superabrasive particles **10** may be about 75 weight percent to about 99 weight percent diamond particles with an average size of about 10 μm to about 40 μm and the balance being diamond particles exhibiting an average size of about 1 μm to about 10 μm . It should be understood that any of the above ranges of various sizes and weight percentages for the coarse superabrasive particles **10** and the ultra-dispersed diamond particles **12** may be combined and processed according to any of the methods disclosed herein (i.e. time, temperature, and pressure parameters). As mentioned above, in one embodiment, superabrasive particles **10** may comprise diamond. In other embodiments, superabrasive particles **10** may comprise boron nitride, silicon carbide, etc., without limitation.

Referring now to both FIGS. 1 and 2, the sintered superabrasive element **2** will exhibit, to a substantial extent, the same or similar grain/particle size as the grain/particle size and relative weight percentages of the precursor materials used (i.e., the coarse superabrasive particles **10** and ultra-dispersed diamond particles **12**). Additionally, some carbon phases present in the ultra-dispersed diamond particles **12** may be present in the final sintered superabrasive element **2** in detectable small amounts. Such phases include small amounts of amorphous carbon, carbon onion, or both that are not transformed to diamond or otherwise transformed during the HPHT sintering process.

In many applications, it may be desirable to form the superabrasive element **2** on a substrate. For example, in one embodiment, the superabrasive element **2** may comprise a PDC cutting element. FIG. 3 shows a schematic illustration of the process for forming any of the superabrasive elements **2** disclosed herein on a substrate to form a PDC cutting element. With reference to FIG. 3, the mixture **8** of the coarse superabrasive particles **10** and ultra-dispersed diamond particles **12** is positioned adjacent to an interfacial surface **16** of a suitable substrate **18**. As shown in FIG. 3, in one embodiment, substrate **18** may be generally cylindrical. More generally, the substrate **18** may comprise any selected shape and size, without limitation. Although FIG. 3 shows the interfacial surface **16** as being substantially planar, the interfacial surface **16** may exhibit a selected nonplanar topography, without limitation. The substrate **18** may comprise, for example, cobalt-cemented tungsten carbide or another suitable material. Other materials that may be used for the substrate **18** include, without limitation, cemented carbides including titanium carbide, niobium carbide, tantalum carbide, vanadium carbide, iron, nickel, and combinations thereof. The mixture **8** and the substrate **14** may be subjected to an HPHT sintering process to form a superabrasive table **13** (e.g., a PCD table) bonded to the interfacial surface **16** of substrate **18** to form an element suitable for use as a cutting element **15** (e.g., a PDC cutting element).

As previously discussed, a catalyst material (e.g., cobalt, nickel, iron, alloys of the foregoing, etc.) may be employed for promoting intergrowth between the coarse superabrasive particles **10**, the ultra-dispersed diamond particles **12**, and/or intergrowth between the coarse superabrasive particles grains **10** and the ultra-dispersed diamond particles **12**. For example, if superabrasive particles **10** comprise diamond, when the mixture **8** is placed adjacent to, for example, a cobalt-cemented tungsten carbide substrate **18** and subjected to the HPHT sintering process, molten cobalt may wick or

sweep into the mixture **8** from the substrate **18**. Such cobalt may remain in the superabrasive (e.g., PCD) table **13** upon sintering and cooling. One specific embodiment of a non-cobalt catalyst is the INVAR® alloy. In other embodiments, a catalyst may be provided prior to sintering, as a layer of material between the substrate **18** and the mixture **8**, within the mixture **8** as a powder, or as otherwise known in the art. Of course, when the superabrasive table **13** includes a catalyst material, the thermal stability may be lower compared to when the superabrasive table **13** is fabricated without the catalyst material. Moreover, such a catalyst material may be at least partially removed (e.g., by acid-leaching or as otherwise known in the art) from at least a portion of the superabrasive table **13** to a selected depth from an exterior surface of the superabrasive table **13**, if desired.

Still, in other embodiments, the superabrasive table **13** may be sintered and, subsequently, bonded to the substrate **18** by brazing, using an HPHT process, or any other suitable joining technique, without limitation.

FIGS. **4** and **5** show an isometric view and a top elevation view, respectively, of a rotary drill bit **20** in accordance with one embodiment of the present invention. The rotary drill bit **20** includes at least one superabrasive cutting element manufactured in accordance with embodiments of the present invention. The rotary drill bit **20** includes a bit body **22** including radially and longitudinally extending blades **38** with leading faces **40**, and a threaded pin connection **50** for connecting the bit body **22** to a drilling string. The bit body **22** defines a leading end structure for drilling into a subterranean formation by rotation about a longitudinal axis **36** and application of weight-on-bit. In one embodiment, at least one superabrasive cutting element comprising polycrystalline diamond may be affixed to rotary drill bit **20**. For example, as best shown in FIG. **5**, a plurality of cutting elements **24** are secured to the blades **38**, each cutting element **24** including a PCD table **28** bonded to a substrate **26**. The PCD tables **28** and substrates **26** may be fabricated and structured in accordance with any of the disclosed embodiments. More generally, the diamond tables **28** may comprise any superabrasive encompassed by the instant disclosure, without limitation. In addition, if desired, in some embodiments, some of the cutting elements **24** may be conventional in construction, if desired. Also, circumferentially adjacent blades **38** define so-called junk slots **42** therebetween, as known in the art. Additionally, the rotary drill bit **20** includes a plurality of nozzle cavities **46** for communicating drilling fluid from the interior of the rotary drill bit **20** to the cutting elements **24**.

FIGS. **4** and **5** merely depict one embodiment of a rotary drill bit that employs at least one cutting element including a superabrasive element fabricated and structured in accordance with the disclosed embodiments, without limitation. The drill bit **20** is used to represent any number of earth-boring tools or drilling tools, including, for example, core bits, roller-cone bits, fixed-cutter bits, eccentric bits, bicenter bits, reamers, reamer wings, or any other downhole tool including superabrasive cutting elements or inserts, without limitation.

The superabrasive elements disclosed herein may also be utilized in applications other than cutting technology. The embodiments of superabrasive elements disclosed herein may be used in wire dies, bearings, artificial joints, inserts, cutting elements, and heat sinks. Thus, any of the superabrasive elements formed by the methods disclosed herein may be employed for forming an article of manufacture including at least one superabrasive element.

Thus, the embodiments of superabrasive elements disclosed herein may be used on any apparatus or structure in which at least one conventional PCD element is typically used. In one embodiment, a rotor and a stator (i.e., a thrust bearing apparatus) may each include a superabrasive element according to any of the embodiments disclosed herein and may be operably assembled to a downhole drilling assembly. U.S. Pat. Nos. 4,410,054; 4,560,014; 5,364,192; 5,368,398; and 5,480,233, the disclosure of each of which is incorporated herein, in its entirety, by this reference, disclose subterranean drilling systems within which bearing apparatuses utilizing PCD elements disclosed herein may be incorporated. The embodiments of superabrasive elements disclosed herein may also form all or part of heat sinks, wire dies, bearing elements, cutting elements, cutting inserts (e.g., on a roller cone type drill bit), machining inserts, or any other article of manufacture as known in the art. Other examples of articles of manufacture that may use any of PCD elements disclosed herein are disclosed in U.S. Pat. Nos. 4,811,801; 4,274,900; 4,268,276; 4,468,138; 4,738,322; 4,913,247; 5,016,718; 5,092,687; 5,120,327; 5,135,061; 5,154,245; 5,460,233; 5,544,713; and 6,793,681, the disclosure of each of which is incorporated herein, in its entirety, by this reference.

The following working examples of the present invention set forth various formulations that have been used to form PDC cutting elements. The following working examples provide further detail in connection with the specific embodiments described above.

COMPARATIVE EXAMPLE 1

A conventional PDC was formed from a mixture of coarse diamond particles being in excess of about 90 weight percent of the mixture with the balance being relatively fine diamond particles. The coarse diamond particles exhibit an average particle size about ten times the average particle size of the relatively fine diamond particles of the mixture. The mixture contained no ultra-dispersed diamond particles. The mixture was placed adjacent to a cobalt-cemented tungsten carbide substrate. The mixture and substrate were placed in a niobium can and sintered at a temperature of about 1400° C. and about 69 kilobar for about 90 seconds to form the conventional PDC. The thermal stability of the as-formed conventional PDC was evaluated by measuring the distance cut in a granite workpiece prior to failure without using coolant. The distance cut is considered representative of the thermal stability of the PDC. The conventional PDC was able to cut a distance of only 1643 linear feet in the granite workpiece.

EXAMPLE 2

A mixture was formed of about 0.1 weight percent ultra-dispersed diamond particles with the balance comprising the same diamond formulation as example 1. The mixture was placed adjacent to a cobalt-cemented tungsten carbide substrate. The mixture and substrate were placed within a niobium can and sintered at a temperature of about 1400° C. and about 69 kilobar for about 90 seconds to form a PDC. The thermal stability of the as-formed PDC was evaluated by measuring the distance cut in a granite workpiece without using coolant. The PDC fabricated in example 2 was able to cut a distance of 2135 linear feet in a granite workpiece compared to the distance of only 1643 linear feet cut by the conventional PDC of example 1. Thus, thermal stability tests indicate that the PDC of example 2 exhibited a significantly

11

improved thermal stability compared to the conventional PDC of comparative example 1. Wear flat volume tests indicated that there was no substantial difference in wear resistance between the PDC of example 2 and the conventional PDC of comparative example 1.

EXAMPLE 3

A mixture was formed of about 1 weight percent ultra-dispersed diamond particles with the balance comprising the same diamond formulation as example 1. The mixture was placed adjacent to a cobalt-cemented tungsten carbide substrate. The mixture and substrate were placed within a niobium can and sintered at a temperature of about 1400° C. and about 69 kilobar for about 90 seconds to form a PDC. The PDC fabricated in example 3 was able to cut a distance of 1868 linear feet in a granite workpiece compared to a distance of only 1643 linear feet cut by the conventional PDC of example 1. Thus, thermal stability tests indicate that the PDC of example 3 also exhibited a significantly improved thermal stability compared to the conventional PDC of example 1. Wear flat volume tests indicated that there was no substantial difference in wear resistance between the PDC of example 3 and the conventional PDC of example 1.

From the foregoing it will be appreciated that, although specific embodiments and working examples of the present invention have been described herein for purposes of illustration, various modifications may be made without deviating from the spirit and scope of the present invention. Accordingly, the present invention is not limited except as by the appended claims. The words "including" and "having," as used herein, including the claims, shall have the same meaning as the word "comprising."

The invention claimed is:

1. A method of forming an abrasive element, comprising: sintering a powder mixture including a plurality of superabrasive particles, a plurality of sp²-carbon-containing particles, and a metallic constituent to form the abrasive element; and wherein the plurality of sp²-carbon-containing particles are present in the powder mixture in an amount greater than zero weight percent to about 5 weight percent of the powder mixture.
2. The method of claim 1 wherein the plurality of superabrasive particles includes at least one of diamond, carbide, or boron nitride.
3. The method of claim 1 wherein the plurality of superabrasive particles includes diamond particles.
4. The method of claim 1 wherein the plurality of superabrasive particles includes carbide.
5. The method of claim 4 wherein the carbide includes silicon carbide.
6. The method of claim 1 wherein the plurality of sp²-carbon-containing particles includes at least one of ultra-dispersed diamond particles or nanocrystalline diamond.

12

7. The method of claim 1 wherein the plurality of sp²-carbon-containing particles are present in the powder mixture in an amount greater than zero weight percent to about 4 weight percent of the powder mixture.

8. The method of claim 1 wherein the plurality of sp²-carbon-containing particles are present in the powder mixture in an amount greater than 0.1 weight percent to about 1.5 weight percent of the powder mixture.

9. The method of claim 1 wherein the plurality of sp²-carbon-containing particles are present in the powder mixture in an amount greater than 0.1 weight percent to about 1 weight percent of the powder mixture.

10. The method of claim 1, further comprising attaching the abrasive element to a substrate.

11. The method of claim 1 wherein the metallic constituent includes iron, nickel, cobalt, or alloys thereof.

12. The method of claim 1 wherein the metallic constituent includes a catalyst material.

13. The method of claim 1 wherein the abrasive element forms a component of a superabrasive cutting element.

14. The method of claim 1 wherein sintering a powder mixture including a plurality of superabrasive particles, a plurality of sp²-carbon-containing particles, and a metallic constituent to form the abrasive element includes subjecting the powder mixture to a high-pressure/high-temperature process.

15. A method of forming a superabrasive element, comprising:

sintering a powder mixture including a plurality of diamond particles, a plurality of sp²-carbon-containing particles, a plurality of carbide particles, and a metallic constituent to form the superabrasive element;

wherein the plurality of sp²-carbon-containing particles are present in the powder mixture in an amount greater than zero weight percent to about 45 weight percent of the powdered mixture.

16. The method of claim 15 wherein the plurality of sp²-carbon-containing particles includes at least one of ultra-dispersed diamond particles or nanocrystalline diamond.

17. The method of claim 15 wherein the plurality of sp²-carbon-containing particles comprises greater than zero weight percent to about 1 weight percent of the powder mixture.

18. The method of claim 15 wherein the plurality of sp²-carbon-containing particles comprises greater than zero weight percent to about 2 weight percent of the powder mixture.

19. The method of claim 15 wherein the plurality of carbide particles includes silicon carbide.

20. The method of claim 15 wherein the metallic constituent includes iron, nickel, cobalt, or alloys thereof.

* * * * *